

[BILLING CODE 4140-01-P]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB review; comment request

The SSA-NIH Collaboration to Improve the Disability Determination Process: Validation of IRT-CAT Tools

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Clinical Research Center, the National Institutes of Health has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the Federal Register, September 8, 2011, Volume 76, Number 174, page 55690, and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

Potential persons who are to respond to the collection of information are not required to respond to the collection of information unless it displays a currently valid OMB control number.

PROPOSED COLLECTION: *Title:* The SSA-NIH Collaboration to Improve the Disability Determination Process: Validation of IRT-CAT tools. Type of Information Collection Reguest: NEW. Need and Use of Information Collection: The Epidemiology and Biostatistics section in RMD will be collecting information through a contractor (Boston University- Health and Disability Research Institute (BU-HDR)) and subcontractor for validation of the Computer Adaptive Tests which are being developed to assist in the SSA disability determination process. The utilization of CAT technology could potentially allow the SSA to collect more relevant and precise data about human functioning in a faster, more efficient fashion. To validate the CAT assessments that have been developed, the contractor will administer both the BU-HDR CAT and established legacy instruments in a small sample of adults who report their current employment status as "permanently disabled". Individuals will complete the CAT tools for the functional domains of Physical Demands and Interpersonal Interactions along with established legacy instruments. For the domain of physical function, individuals will complete the BU-HDR CAT; the PROMIS Item Bank v 1.0-Physical Functioning © PROMIS Health Organization and PROMIS Cooperative Group;

and, The Short Form (36) Health Survey™ (SF-36). For the domain of interpersonal interactions, individuals will complete the BU-HDR CAT, the SF-36 and the BASIS-24© (Behavior and Symptom Identification Scale). Data collected will be used to validate the BU-HDR CAT tools. Without this information, completion of the BU-HDR CAT tools will not be possible. Frequency of Response: Once. Affected Public: Individuals who have opted in to participate in web surveys through a survey research firm. Type of Respondents: Adults who indicate "permanently disabled" as a working status. The annual reporting burden is as follows:

ESTIMATES OF HOUR BURDEN				
Type of	Number of	Frequency	Average	Annual Hour
Respondents	Respondents	of	Time per	Burden
		Response	Response	
Patients	1,000	1	0.5	500.00
Totals				500.00

REQUEST FOR COMMENTS: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

DIRECT COMMENTS TO OMB: Written comments and/or suggestions regarding the item(s) contained in this

notice, especially regarding the estimated public burden and associated response time, should be directed to the:

Office of Management and Budget, Office of Regulatory Affairs, OIRA_submission@omb.eop.gov or by fax to

202-395-6974, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain

a copy of the data collection plans and instruments, contact: Ms. Meghan Gleason, Rehabilitation Medicine

Department, Clinical Research Center, NIH, Building 10, Room 1-2420, 9000 Rockville Pike, Bethesda, MD

20892, or call non-toll-free number (301) 443-9085 or E-mail your request, including your address to:

meghan.gleason@nih.gov

COMMENTS DUE DATE: Comments regarding this information collection are best assured of having their full

effect if received within 30-days of the date of this publication.

Date: December 5, 2011

Elizabeth K Rasch,

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Clinical Research Center

National Institutes of Health

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